

07-03-00

A

ATTORNEY DOCKET NO. 21011.0038U1

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

BOX PATENT APPLICATION
Assistant Commissioner for Patents
Washington, D.C. 20231

NEEDLE & ROSENBERG, P.C.
Suite 1200, The Candler Building
127 Peachtree Street, N.E.
Atlanta, Georgia 30303-1811

June 30, 2000

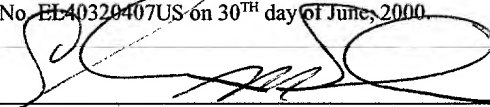
Dear Sir:

Transmitted herewith for filing are the specification and claims of the patent application of:

Inventor(s): Michael Joseph Banner; Paul Bradford Blanch; Neil Russell Euliano, II
and Jose C. Principe

Title of Invention: "VENTILATOR MONITOR SYSTEM AND METHOD OF USING
SAME"

Also enclosed are:

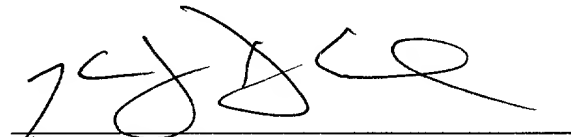
9	SHEETS OF	<input type="checkbox"/> FORMAL DRAWINGS	<input checked="" type="checkbox"/> INFORMAL DRAWINGS
	OATH OR DECLARATION OF APPLICANT(S)		
	A POWER OF ATTORNEY		
	A PRELIMINARY AMENDMENT		
X	A VERIFIED STATEMENT TO ESTABLISH SMALL ENTITY STATUS UNDER 37 C.F.R. §1.9 AND §1.27		
X	A CHECK IN THE AMOUNT OF \$546.00 TO COVER THE FILING FEE.		
X	THE COMMISSIONER IS HEREBY AUTHORIZED TO CHARGE ANY ADDITIONAL FEES WHICH MAY BE REQUIRED IN CONNECTION WITH THE FOLLOWING OR CREDIT ANY OVERPAYMENT TO ACCOUNT NO. 14-0629		
	A CERTIFIED COPY OF PREVIOUSLY FILED FOREIGN APPLICATION NO. FILED IN ON .		
X	I hereby certify that this correspondence is being placed in the United States Mail as Express Mail No. EL40320407US on 30 TH day of June, 2000.  Everardo McFarlane 6-30-2000 DATE		
	A computer readable form of the sequence listing in compliance with 37 C.F.R. § 1.821(e). The content of the computer readable form of the sequence listing and the sequence listing in the specification of the application as filed are the same.		
	OTHER (IDENTIFY)		

The filing fee is calculated as follows:

CLAIMS AS FILED, LESS ANY CLAIMS CANCELLED BY AMENDMENT

TOTAL CLAIMS = $38 - 20 = 18 \times \$18.00 =$	\$324.00
INDEPENDENT CLAIMS = $3 - 3 = 0 \times \$78.00 =$	\$78.00
BASIC FEE =	\$690.00
TOTAL OF ABOVE CALCULATIONS =	\$1,092.00
REDUCTION BY 1/2 FOR SMALL ENTITY =	\$546.00
TOTAL FILING FEE =	\$546.00

Respectfully submitted,



Kean J. DeCarlo

Registration No. 39,954

NEEDLE & ROSENBERG, P.C.
Suite 1200, The Candler Building
127 Peachtree Street, N.E.
Atlanta, Georgia 30303-1811
(404) 688-0770

APPLICANTS OR PATENTEES: Michael J. Banner; Paul Bradford Blanch and
Neil Russell Euliano II

FOR: "VENTILATOR MONITOR SYSTEM AND METHOD
OF USING SAME"

**VERIFIED STATEMENT (DECLARATION) CLAIMING SMALL ENTITY
STATUS (37 CFR 1.9(f) and 1.27(d)) - NONPROFIT ORGANIZATION**

I hereby declare that I am an official empowered to act on behalf of the nonprofit organization
identified below:

NAME OF ORGANIZATION: THE UNIVERSITY OF FLORIDA
ADDRESS OF ORGANIZATION: 223 Grinter Hall
Gainesville, Florida 32611-5500

TYPE OF ORGANIZATION:

- ☒ University or other institution of higher education
(Name of state Florida)
(Citation of statute)
☐ Tax exempt under Internal Revenue Service Code (26 USC 501(a) and 501(c)(3))
☐ Non-profit scientific or educational under statute of state of the United States of
America
(Name of state)
(Citation of statute)
☐ Would qualify as tax exempt under Internal Revenue Service Code (26 USC 501(a)
and 501(c) (3)) if it were located in The United States of America
☐ Would qualify as nonprofit scientific or educational under statute of state of The
United States of America if it were located in The United States of America
(Name of state)
(Citation of statute)

I hereby declare that the nonprofit organization identified above qualifies as a nonprofit
organization as defined in 37 CFR 1.9 (e) for purposes of paying reduced fees under section 41(a)
and (b) of Title 35, United States Code with regard to the invention entitled "**VENTILATOR
MONITOR SYSTEM AND METHOD OF USING SAME**" by inventors Michael J. Banner,
Paul Bradford Blanch, Neil Russell Euliano, II, and Jose C. Principe described in the specification
filed herewith.

I hereby declare that rights under contract or law have been conveyed to and remain with the
nonprofit organization with regard to the above identified invention.

If the rights held by the nonprofit organization are not exclusive, each individual, concern or
organization having rights to the invention is listed below* and no rights to the invention are held
by any person, other than the inventor, who could not qualify as a small business concern under 37
CFR 1.9(d) or by any concern which would not qualify as a small business concern under 37 CFR
1.9(d) or a nonprofit organization under 37 CFR 1.9(e).

ATTORNEY DOCKET NO. 21011.0038U1

*NOTE: Separate verified statements are required for each named person, concern or organization having rights to the invention averring to their status as small entities. (37 CFR 1.27)

NAME _____
ADDRESS _____

☐ INDIVIDUAL ☐ SMALL BUSINESS CONCERN ☐ NONPROFIT ORGANIZATION

I acknowledge the duty to file, in this application or patent, notification of any change in status resulting in loss of entitlement to small entity status prior to paying, or at the time of paying, the earliest of the issue fee or any maintenance fee due after the date on which status as a small entity is no longer appropriate. (37 CFR 1.28(b)).

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon or any patent to which this verified statement is directed.

NAME OF PERSON SIGNING: Winfred M. Phillips

TITLE IN ORGANIZATION: Vice President of Research & Graduate Programs

ADDRESS OF PERSON SIGNING: 223 Grinter Hall, Gainesville, FL 32611-5500

SIGNATURE: Winfred M. Phillips

DATE: June 29, 2000

5

Express Mail No. EL403204074US

Attorney Docket No. 21011.0038U1

UTILITY PATENT FILING

10

APPLICATION

15

FOR

UNITED STATES LETTERS PATENT

VENTILATOR MONITOR SYSTEM AND METHOD OF USING SAME

20

BY

25

Michael J. Banner, a citizen of the United States of America, residing at 14727 NW 60th Avenue, Alachua, Florida 32615, **Paul Bradford Blanch**, a citizen of the United States of America, residing at 15214 NW 94th Avenue, Alachua, Florida 32615, **Neil Russell Euliano II**, a citizen of the United States of America, residing at 3914 South West 95th Drive, Gainesville, Florida 32608, and **Jose C. Principe**, a citizen of the United States of America, residing at 3022 North West 24th Terrace, Gainesville, Florida 32605..

30

5
VENTILATOR MONITOR SYSTEM AND METHOD OF USING SAME

10
BACKGROUND OF THE INVENTION

15
Field of the Invention:

The present invention relates to the respiratory care of a patient and, more particularly, to a ventilator monitor system that receives a plurality of ventilator support signals indicative of the sufficiency of ventilation support received by the patient, receives at least one ventilator signal indicative of the level settings of the ventilator setting controls of the ventilator, and determines the desired level settings of the ventilator setting controls of the ventilator to provide the appropriate quality and quantity of ventilation support to the patient.

20
Background:

Mechanical ventilatory support is widely accepted as an effective form of therapy and means for treating patients with respiratory failure. Ventilation is the process of delivering oxygen to and washing carbon dioxide from the alveoli in the lungs. When receiving ventilatory support, the patient becomes part of a complex interactive system which is expected to provide adequate ventilation and promote gas exchange to aid in the stabilization and recovery of the patient. Clinical treatment of a ventilated patient often calls for monitoring a patient's breathing to detect an interruption or an irregularity in the breathing pattern, for triggering a ventilator to initiate assisted breathing, and for interrupting the assisted breathing periodically to wean the patient off of the assisted breathing regime, thereby restoring the patient's ability to breath independently.

25
30
In those instances in which a patient requires mechanical ventilation due to respiratory failure, a wide variety of mechanical ventilators are available. Most

5 modern ventilators allow the clinician to select and use several modes of inhalation
either individually or in combination via the ventilator setting controls that are
common to the ventilators. These modes can be defined in three broad categories:
spontaneous, assisted or controlled. During spontaneous ventilation without other
10 modes of ventilation, the patient breathes at his own pace, but other interventions may
affect other parameters of ventilation including the tidal volume and the baseline
pressure, above ambient, within the system. In assisted ventilation, the patient
initiates the inhalation by lowering the baseline pressure by varying degrees, and then
the ventilator "assists" the patient by completing the breath by the application of
positive pressure. During controlled ventilation, the patient is unable to breathe
15 spontaneously or initiate a breath, and is therefore dependent on the ventilator for
every breath. During spontaneous or assisted ventilation, the patient is required to
"work" (to varying degrees) by using the respiratory muscles in order to breath.

20 The work of breathing (the work to initiate and sustain a breath) performed by
a patient to inhale while intubated and attached to the ventilator may be divided into
two major components: physiologic work of breathing (the work of breathing of the
patient) and breathing apparatus imposed resistive work of breathing. The work of
breathing can be measured and quantified in Joules/L of ventilation. In the past,
techniques have been devised to supply ventilatory therapy to patients for the purpose
25 of improving patient's efforts to breath by decreasing the work of breathing to sustain
the breath. Still other techniques have been developed that aid in the reduction of the
patient's inspiratory work required to trigger a ventilator system "ON" to assist the
patient's breathing. It is desirable to reduce the effort expended by the patient in each
of these phases, since a high work of breathing load can cause further damage to a
30 weakened patient or be beyond the capacity or capability of small or disabled patients.
It is further desirable to deliver the most appropriate mode, and, intra-mode, the most
appropriate quality and quantity of ventilation support required the patient's current
physiological needs.

5 The early generation of mechanical ventilators, prior to the mid-1960s, were
designed to support alveolar ventilation and to provide supplemental oxygen for those
patients who were unable to breathe due to neuromuscular impairment. Since that
time, mechanical ventilators have become more sophisticated and complicated in
10 response to increasing understanding of lung pathophysiology. Larger tidal volumes,
an occasional "sigh breath," and a low level of positive end-expiratory pressure
(PEEP) were introduced to overcome the gradual decrease in functional residual
capacity (FRC) that occurs during positive-pressure ventilation (PPV) with lower tidal
volumes and no PEEP. Because a decreased functional residual capacity is the
15 primary pulmonary defect during acute lung injury, continuous positive pressure
(CPAP) and PEEP became the primary modes of ventilatory support during acute
lung injury.

20 In an effort to improve a patient's tolerance of mechanical ventilation, assisted
or patient-triggered ventilation modes were developed. Partial PPV support, in which
mechanical support supplements spontaneous ventilation, became possible for adults
outside the operating room when intermittent mandatory ventilation (IMV) became
available in the 1970s. Varieties of "alternative" ventilation modes addressing the
needs of severely impaired patients continue to be developed.

25 The second generation of ventilators was characterized by better electronics
but, unfortunately, due to attempts to replace the continuous high gas flow IMV
system with imperfect demand flow valves, failed to deliver high flow rates of gas in
response to the patient's inspiratory effort. This apparent advance forced patient to
perform excessive imposed work and thus, total work in order to overcome ventilator,
30 circuit, and demand flow valve resistance and inertia. In recent years,
microprocessors have been introduced into modern ventilators. Microprocessor
ventilators are typically equipped with sensors that monitor breath-by-breath flow,
pressure, volume, and derive mechanical respiratory parameters. Their ability to
sense and transduce "accurately," combined with computer technology, makes the

5 interaction between clinician, patient, and ventilator more sophisticated than ever.
The prior art microprocessor controlled ventilators suffered from compromised
accuracy due to the placement of the sensors required to transduce the data signals.
Consequently, complicated algorithms were developed so that the ventilators could
"approximate" what was actually occurring within the patient's lungs on a breath by
10 breath basis. In effect, the computer controlled prior art ventilators were limited to
the precise, and unyielding, nature of the mathematical algorithms which attempted to
mimic cause and effect in the ventilator support provided to the patient.

15 Unfortunately, as ventilators become more complicated and offer more
options, the number of potentially dangerous clinical decisions increases. The
physicians, nurses, and respiratory therapists that care for the critically ill are faced
with expensive, complicated machines with few clear guidelines for their effective
use. The setting, monitoring, and interpretation of some ventilatory parameters have
become more speculative and empirical, leading to potentially hazardous misuse of
20 these new ventilator modalities. For example, the physician taking care of the patient
may decide to increase the pressure support ventilation (PSV) level based on the
displayed spontaneous breathing frequency. This may result in an increase in the
work of breathing of the patient which may not be appropriate. This "parameter-
monitor" approach, unfortunately, threatens the patient with the provision of
25 inappropriate levels of pressure support.

30 Ideally, ventilatory support should be tailored to each patient's existing
pathophysiology, rather than employing a single technique for all patients with
ventilatory failure (*i.e.*, in the example above, of the fallacy of using spontaneous
breathing frequency to accurately infer a patient's work of breathing). Thus, current
ventilatory support ranges from controlled mechanical ventilation to total spontaneous
ventilation with CPAP for support of oxygenation and the elastic work of breathing
and restoration of lung volume. Partial ventilation support bridges the gap for patients
who are able to provide some ventilation effort but who cannot entirely support their

5 own alveolar ventilation. The decision-making process regarding the quality and
quantity of ventilatory support is further complicated by the increasing knowledge of
the effect of mechanical ventilation on other organ systems.

10 The overall performance of the assisted ventilatory system is determined by
both physiological and mechanical factors. The physiological determinants, which
include the nature of the pulmonary disease, the ventilatory efforts of the patient, and
many other physiological variables, changes with time and are difficult to diagnosis.
Moreover, the physician historically had relatively little control over these
15 determinants. Mechanical input to the system, on the other hand, is to a large extent
controlled and can be reasonably well characterized by examining the parameters of
ventilator flow, volume, and/or pressure. Optimal ventilatory assistance requires both
appropriately minimizing physiologic workloads to a tolerable level and decreasing
imposed resistive workloads to zero. Doing both should insure that the patient is
neither overstressed nor oversupported. Insufficient ventilatory support places
20 unnecessary demands upon the patient's already compromised respiratory system,
thereby inducing or increasing respiratory muscle fatigue. Excessive ventilatory
support places the patient at risk for pulmonary-barotrauma, respiratory muscle
deconditioning, and other complications of mechanical ventilation.

25 Unfortunately, none of the techniques devised to supply ventilatory support
for the purpose of improving patient efforts to breath, by automatically decreasing
imposed work of breathing to zero and appropriately decreasing physiologic work
once a ventilator system has been triggered by a patient's inspiratory effort, provides
the clinician with advice in the increasingly complicated decision-making process
30 regarding the quality and quantity of ventilatory support. As noted above, it is
desirable to reduce the effort expended by the patient to avoid unnecessary medical
complications of the required respiratory support and to deliver the most appropriate
mode, and, intra-mode, the most appropriate quality and quantity of ventilation
support required the patient's current physiological needs. Even using the advanced

5 microprocessor controlled modern ventilators, the prior art apparatus and methods
tend to depend upon mathematical models for determination of necessary actions. For
example, a ventilator may sense that the hemoglobin oxygen saturation level of the
patient is inappropriately low and, from the sensed data and based upon a determined
10 mathematical relationship, the ventilator may determine that the oxygen content of the
breathing gas supplied to the patient should be increased. This is similar to, and
unfortunately as inaccurate as, a physician simply looking at a patient turning “blue”
and determining more oxygen is needed.

15 From the above, in the complicated decision-making environment engendered
by the modern ventilator, it is clear that it would be desirable to have a medical
ventilator monitor system that alerts the clinician of the ventilator’s failure to supply
the appropriate quality and quantity of ventilatory support and provides advice to the
clinician regarding the appropriate quality and quantity of ventilatory support that is
20 tailored to the patient’s pathophysiology. Such a ventilatory monitor system is
unavailable in current systems.

SUMMARY

25 In accordance with the purposes of this invention, as embodied and broadly
described herein, this invention, in one aspect, relates to a method of monitoring the
ventilation support provided by a ventilator that is supplying a breathing gas to a
patient via a breathing circuit that is in fluid communication with the lungs of the
patient. The ventilator has a plurality of selectable ventilator setting controls
30 governing the supply of ventilation support from the ventilator, each setting control
selectable to a level setting. The ventilator support monitor system preferably
receives at least one ventilator setting parameter signal, each ventilator setting
parameter signal indicative of the level settings of one ventilator setting control,
monitors a plurality of sensors, each sensor producing an output signal indicative of a

5 measured ventilation support parameter, to determine the sufficiency of the ventilation
support received by the patient, and determines the desired level settings of the
ventilator setting controls in response to the received ventilator setting parameter
signal and the output signals. The ventilator support monitor system preferably
utilizes a trainable neural network to determine the desired level settings of the
10 ventilator setting controls.

In another aspect, the invention relates to a ventilator support monitor system
that supplies a breathing gas to a patient via a breathing circuit in fluid
communication with the ventilator and the lungs of a patient. The ventilator
preferably has at least one selectable ventilator setting control. The selectable
15 ventilator setting control governs the supply of ventilation support from the ventilator
to the patient via the breathing circuit. Each ventilator setting control generates a
ventilator setting parameter signal indicative of the current level setting of the
ventilator setting.

The ventilator support monitor system has a plurality of sensors and a
processing subsystem. The sensors measure a plurality of ventilation support
parameters and each sensor generates an output signal based on the measured
ventilation support parameter. The processing subsystem is connected to receive the
25 output signal from the sensor and the ventilator setting signal(s) from the ventilator
setting control(s). The processor of the processing subsystem runs under control of a
program stored in the memory of the processing subsystem and determines a desired
level setting of at least one ventilator setting control in response to the ventilator
setting parameter signal and the output signal. The processing subsystem of the
30 ventilator preferably utilizes a trainable neural network to determine the desired level
settings of the ventilator setting controls.

5

DETAILED DESCRIPTION OF THE FIGURES

The accompanying drawings, which are incorporated in and constitute a part of this specification, illustrate several embodiments of the invention and together with the description, serve to explain the principals of the invention.

10

Fig. 1 is a block diagram of one configuration a ventilator monitor system for determining the desired ventilator control settings of a ventilator.

15

Fig. 2A is a block diagram of one configuration of the ventilator monitor system showing the ventilator providing ventilation support to a patient connected to the ventilator via a breathing circuit.

20

Fig. 2B is a block diagram of an embodiment of a ventilator monitor system showing the monitor system incorporated into the ventilator.

25

Fig. 3 is a block diagram of the ventilator monitor system showing a plurality of sensors connected to the processing subsystem.

Fig. 4 is a block diagram of a processing subsystem of the present invention.

Fig. 5 is a block diagram of a feature extraction subsystem of the present invention.

30

Fig. 6A is a block diagram of one embodiment of the intelligence subsystem of the processing subsystem.

Fig. 6B is a block diagram of a second embodiment of the intelligence subsystem of the processing subsystem.

5 Fig. 7 is a schematic block diagram of one realization of the system of the invention.

 Fig. 8 is a diagram of the basic structure of an artificial neural network having a layered structure.

10

 DETAILED DESCRIPTION OF THE INVENTION

 The present invention is more particularly described in the following examples
15 that are intended to be illustrative only since numerous modifications and variations therein will be apparent to those skilled in the art. As used in the specification and in the claims, the singular form "a," "an" and "the" include plural referents unless the context clearly dictates otherwise.

20 As depicted in Figs. 1 - 3, the ventilator monitor system 10 of the present invention preferably comprises a conventional ventilator 20, a processing subsystem 40, a measuring system, and a display 62. The ventilator 20 supplies a breathing gas to the lungs of the patient P via a breathing circuit 22 that typically comprises an inspiratory line 23, an expiratory line 24, and a patient connection tube 25, all
25 connected by a patient connector 26. The preferred ventilator 20 is a microprocessor-controlled ventilator of a type that is exemplified by a Mallinckrodt, Nelcor, Puritan-Bennet, 7200ae, or a Bird 6400 Ventilator.

30 To control the delivery of the breathing gas, the preferred ventilator 20 typically has at least one selectable ventilator setting control 30 operatively connected to the processing system 40 for governing the supply of ventilation support provided to the patient P. As one skilled in the art will appreciate, each ventilator setting control 30 is selectable to a desired level setting. Such a ventilator 20 is particularly

5 useful in controlling the delivery of breathing support so that the quantity and quality of ventilation support coincides with the physiological support needs of the patient P.

10 In the preferred embodiment, the preferred ventilator 20 can operate selectively in one or more conventional modes, as needed and selected by the operator and/or the processing subsystem 40, including but not limited to: (i) assist control ventilation (ACMV); (ii) synchronized intermittent mandatory ventilation (SIMV); (iii) continuous positive airway pressure (CPAP); (iv) pressure-control ventilation (PCV); (v) pressure support ventilation (PSV); (vi) proportional assist ventilation (PAV); and (vii) volume assured pressure support (VAPS). Further, the level setting of one or more conventional ventilator setting controls 30 of the ventilator 20 (*i.e.*, the intra-mode setting controls of the ventilator 20) may be adjusted, as needed and selected by the operator and/or the processing system 40 in order to maintain the sufficiency of ventilation support delivered to the patient P. The ventilator setting controls 30 of the ventilator 20 include but are not limited to controls for setting: (i) a minute ventilation (V_e) level; (ii) a ventilator breathing frequency (f) level; (iii) a tidal volume (V_T) level; (iv) a breathing gas flow rate (V) level; (v) a pressure limit level; (vi) a work of breathing (WOB) level; (vii) a pressure support ventilation (PSV) level; (viii) a positive end expiratory pressure (PEEP) level; (ix) a continuous positive airway pressure (CPAP) level; and (x) a fractional inhaled oxygen concentration (FIO_2) level.

25 The conventional ventilator 20 contemplated typically has a gas delivery system and may also have a gas composition control system. The gas delivery system may, for example, be a pneumatic subsystem 32 in fluid/flow communication with a gas source 34 of one or more breathing gases and the breathing circuit 22 and in operative connection with the ventilator control settings 30 of the ventilator 20 and the processing subsystem 40. The breathing circuit 22 is in fluid communication with the lungs of the patient P. As one skilled in the art will appreciate, the pneumatic subsystem 40 of the ventilator 20 and the operative connection of that pneumatic

5 subsystem 40 to the source of breathing gas 34 of the ventilator 20 may be any design known in the art that has at least one actuator (not shown) that is capable of being operatively coupled, preferably electrically coupled, to the ventilator setting controls 30 for control of, for example, the flow rate, frequency, and/or pressure of the breathing gas delivered by the ventilator 20 to the patient P from the gas source 34. 10 Such a pneumatic system 32 is disclosed in U.S. Patents Nos. 4,838,259 to Gluck *et al.*, 5,303,698 to Tobia *et al.*, 5,400,777 to Olsson *et al.*, 5,429,123 to Shaffer *et al.*, and 5,692,497 to Schnitzer *et al.*, all of which are incorporated in their entirety by reference herein and is exemplified by the Mallinckrodt, Nelcor, Puritan-Bennet, 7200ae, and the Bird 6400 Ventilator.

15 The gas composition control system may, for example, be an oxygen control subsystem 36 coupled to the source of breathing gas 34 and in operative connection to the ventilator setting controls 30 of the ventilator 20 and the processing subsystem 40. The oxygen control subsystem 36 allows for the preferred control of the percentage composition of the gases supplied to the patient P. As one skilled in the art will appreciate, the oxygen control subsystem 36 of the ventilator 20 and the operative connection of that oxygen control subsystem 36 to the pneumatic subsystem 32 and to the source of breathing gas 34 of the ventilator 20 may be any design known in the art that has at least one actuator (not shown) that is capable of being operatively coupled, 20 preferably electrically coupled, to the ventilator setting controls 30 for control of, for example, the percentage composition of the oxygen supplied to the patient P. 25

30 The processing subsystem 40 of the ventilator monitor system 10 preferably has an input 44 that is operatively coupled to the ventilator setting controls 30 of the ventilator 20 so that at least one ventilator setting parameter signal 42 may be received by the processing subsystem 40. Each ventilator setting parameter signal 42 is preferably indicative of a setting of a ventilator setting control 30. Thus, the processing system 40 is in receipt of signals 42, preferably continuously, indicative of the current level settings of the ventilator setting controls 30. As one skilled in the art

5 will appreciate, the current level settings of the ventilator setting controls 30 may be stored in the memory of the processing subsystem 40. In this example, the ventilator setting parameter signals 42 would be input from the memory of the processing subsystem 40 to the processor for continued processing and assessment.

10 For example, the input of the processing system 40 may receive one or more of the following ventilator setting parameter signals 42: a minute ventilation (V_E) signal indicative of the V_E level set on the ventilator 20; a ventilator breathing frequency (f) signal indicative of the f level set on the ventilator 20; a tidal volume (V_T) signal indicative of the V_T level set on the ventilator 20; a breathing gas flow rate (V) signal indicative of the V level set on the ventilator 20; a pressure limit signal indicative of the pressure limit set on the ventilator 20; a work of breathing (WOB) signal indicative of the WOB level set on the ventilator 20; a pressure support ventilation (PSV) signal indicative of the PSV level set on the ventilator 20; a positive end expiratory pressure (PEEP) signal indicative of the PEEP level set on the ventilator 20; a continuous positive airway pressure (CPAP) signal indicative of the CPAP level set on the ventilator 20; and a fractional inhaled oxygen concentration (FIO_2) signal indicative of the FIO_2 level set on the ventilator 20.

25 The measuring system of the monitor system 10 is also operatively connected to the processing subsystem 40. The measuring system senses and measures a plurality of ventilation support parameters which are indicative of the ventilation support provided to the patient P and the physiological condition of the patient P. It is contemplated that the measuring system may comprise at least one sensor 52, and preferably comprises a plurality of sensors 52, for capturing the desired ventilation support data. Each sensor 52 generates an output signal 51 based on the particular measured ventilation support parameter.

30 In one preferred embodiment shown in Fig. 3, the processing subsystem 30 is shown operatively connected to a flow rate sensor 53, a exhaled CO₂ (Ex CO₂)

5 sensor 54, a pressure sensor 55, a blood pressure sensor 56, and a SPO2 sensor 57. In
this embodiment, it is preferred that the monitor system 10 be responsive to the output
signals 51 input into the processing subsystem 40 from, for example: i) the flow rate
10 sensor 53 which is indicative of the flow rate ventilation support parameter of the gas
expired/inspired by the patient P within the breathing circuit 22, ii) the gas pressure
sensor 55 which is indicative of the pressure ventilation support parameter of the
breathing gas within the breathing circuit 22, and iii) the Ex CO2 sensor 54 which is
indicative of the exhaled carbon dioxide ventilation support parameter present in the
15 exhaled gas expired by the patient P within the breathing circuit 22 (*i.e.*, the flow rate
output signal 51 generated by the flow rate sensor 53, the gas pressure output signal
51 generated by the gas pressure sensor 55, and the Ex CO2 output signal 51
generated by the Ex CO2 sensor 54). Optionally, the monitor system 10 may be
responsive to output signals 51 input into the processing subsystem 40 from the
20 output of the blood pressure sensor 56, which is indicative of the blood pressure
ventilation support parameter of the patient P, for example the arterial systolic,
diastolic, and mean blood pressure of the patient P, and the SPO2 sensor 57 which is
indicative of the hemoglobin oxygen saturation level ventilation support parameter of
the patient P (*i.e.*, the blood pressure output signal 51 generated by the blood pressure
25 sensor 56 and the SPO2 output signal 51 generated by the SPO2 sensor 57).

25 The flow rate sensor 53, the pressure sensor 55, and the Ex CO2 sensor 54 are
preferably positioned between the patient connector 26 and the patient connection
tube 25. Alternatively, it is preferred that the pressure sensor 55 be located at the
tracheal end of the patient connection tube 25. The flow rate, pressure, and Ex CO2
30 sensors 53, 55, 54 are exemplified by Novametrics, CO₂SMO+ monitor (which has a
flow rate, pressure and Ex CO2 sensors). The blood pressure sensor 56 and the SPO2
sensor 57 are exemplified by Dynamap, Inc.'s blood pressure sensor and
Novametrics, CO₂SMO+ monitor's SPO2 sensor. The blood pressure sensor 56 and
the SPO2 sensor 57 may be attached to a portion of the patient's body to render the
requisite measurements. For example, the blood pressure sensor 56, here for example

5 shown as a blood pressure cuff, is shown attached to the arm of the patient P and the SPO2 sensor 57, which may, for example, be a pulse oximeter, is shown attached to a finger of the patient 12. One skilled in the art will appreciate, the blood pressure data may be derived from the SPO2 sensor 57 which eliminates the need for the blood pressure sensor 56.

10 Additional standard equipment can include an operator interface 60, which in the preferred embodiment is a membrane keypad, a keyboard, a mouse, or other suitable input device, for providing user inputs of both data and control commands needed to execute the software which implements the various functions of the invention. The operator of the ventilator monitor system 10 of the present invention may provide the processing subsystem 40, via an operator input signal generated by the operator interface 60, with any number of applicable input parameters, such as patient identification information, patient age, patient weight, or other desired patient statistics. It is preferred that the operator input predetermined patient reference data, such as the arterial blood gas ph, the arterial blood gas PaO2, and/or the arterial blood gas PaCO2 of the patient's blood, and/or patient's temperature into the processing subsystem 40 as operator input signals 61 via the operator interface 60. The monitor system 10 may also be responsive to the core body temperature of the patient P which may be input into the processing subsystem 40 as an output signal 51 from a temperature sensor 58 attached to the patient P or as an operator input signal 61 via the operator interface 60.

30 The processing subsystem 40 preferably comprises a processor 46, for example a microprocessor, a hybrid hardware/software system, controller, or computer, and a memory. The output signals 51 and the ventilation data 72 derived from the output signals 51 are stored in the memory of the processing subsystem 40 at user-defined rates, which may be continuous, for as-needed retrieval and analysis. The ventilator setting signal 42 may also be stored in the memory at a user-defined rate. As one skilled with the art will appreciate, any generated signal may be stored in

5 the memory at user-defined rates. The memory may be, for example, a floppy disk drive, a CD drive, internal RAM or hard drive of the associated processor 12.

10 The processing subsystem 40 is responsive to the output signals 51 of the measuring means, the ventilator setting parameter signal(s) 42, and, if provided, the operator input signals 61. The processor 46 runs under the control of a program stored in the memory and has intelligent programming for the determination of at least one desired level setting of the ventilator setting controls 30 based on at least a portion of the output signal 51 from the measuring means, at least a portion of the ventilator setting parameter signal(s) 42 received at the input 44 of the processing subsystem 40, and, if provided, at least a portion of the operator input signals 61.

15 The desired level settings for the ventilator setting controls 30 of the ventilator 20 may include at least one of the group of: i) a minute ventilation (V_E) level indicative of the desired V_E level to set on the ventilator 20; ii) a ventilator breathing frequency (f) level indicative of the desired f level to set on the ventilator 20; iii) a tidal volume (V_T) level indicative of the V_T level to set on the ventilator 20; iv) a breathing gas flow rate (V) level indicative of the V level to set on the ventilator 20; v) a pressure limit level indicative of the pressure limit level to set on the ventilator 20; vi) a work of breathing (WOB) level indicative of the WOB level to set on the ventilator 20; vii) a pressure support ventilation (PSV) level indicative of the PSV level to set on the ventilator 20; viii) a positive end expiratory pressure (PEEP) level indicative of the PEEP level to set on the ventilator 20; ix) a continuous positive airway pressure (CPAP) level indicative of the CPAP level to set on the ventilator 20; and x) a fractional inhaled oxygen concentration (FIO₂) level indicative of the FIO₂ level to set on the ventilator 20.

25 The desired level setting of the ventilator setting controls 30 determined by the processing system 40 of the monitor system 10 may be displayed to the operator via the display. The display of the monitor system 10 preferably comprises a visual

5 display 62 or CRT, electronically coupled to the processing subsystem 40 for
outputting and displaying output display signals generated from the processing
subsystem 40.

10 Still further, the monitor system 10 may have an alarm 21 for alerting the
operator of either a failure of the monitor system 10, such as a power failure or loss of
signal data input, or an inappropriate setting of a ventilator control 30, such as a level
setting of a ventilator setting control 30 currently controlling the delivery of ventilator
support to the patient P differing from a recommended desired level setting of the
15 ventilator setting control 30. Preferably, the alarm 21 comprises a visual and/or audio
alarm, but any means for alerting the operating clinician know to one skilled in the art
may be used. Of course, it is desired to use a backup power supply, such as a battery.

20 Referring to Figs. 4 and 5, the processing subsystem of the preferred
embodiment of the present invention has a means for determining the desired
ventilation control settings 30 of the ventilator 20. The determining means preferably
comprises a feature extraction subsystem 70 and an intelligence subsystem 80. The
feature extraction subsystem 70 has a means for extracting and compiling pertinent
ventilation data features from the input of the measuring means (*i.e.*, the output
signals 51). In effect, the feature extraction subsystem 70 acts as a preprocessor for
25 the intelligence subsystem 80. An example of the feature extraction subsystem 70 is
shown in Fig. 5. Here, a flow rate sensor 53, a gas pressure sensor 55, a SPO2 sensor
57, an Ex CO2 sensor 54, a temperature (T) sensor 58, a blood pressure (BP) sensor
56, of a type described above, and any other desired sensor are operatively connected
to the feature extraction subsystem 70 of the processing subsystem 40. Preferably, the
30 flow rate sensor 53, the gas pressure sensor 55, and the Ex CO2 sensor 54 provide the
only inputs to the monitor system. The other sensor inputs, and the user input, may be
included to increase the reliability and confidence of the determined desired level
settings of the controls 30. The monitor system 10 preferably adjusts the extraction of
ventilator data 72 as a function of the presence or absence of these optional inputs.

5 By making the number of inputs optional, which also makes the required number of sensors 52 comprising the measuring system optional, the number of environments in which the ventilator monitor system 10 can be used is increased.

10 The purpose of the feature extraction subsystem 70 is to calculate and/or identify and extract important variables or features from the output signals 51 produced by the measuring means. For example, from the exemplified required inputs to the feature extraction subsystem 70, *i.e.*, the gas pressure output signal 51, the flow rate output signal 51, and the Ex CO2 output signal 51, a plurality of ventilation data 72 may be derived. The derived ventilation data 72 may comprise:

15 the values of any output signals 51 used, such as, for example, the gas pressure output signal 51, the flow rate output signal 51, and the Ex CO2 output signal 51 output signals 51; the peak inflation pressure (PIP), which is the maximal pressure generated during mechanical ventilation of the lungs; the mean airway pressure (PAW), which is the average positive pressure measured at the airway opening in the patient connection tube 25 or in the breathing circuit 22 over one minute; the positive end

20 expiratory pressure (PEEP), which is the baseline or starting positive pressure prior to mechanical inflation or the positive pressure applied continuously during inhalation and exhalation during spontaneous ventilation; breathing frequency (f), which is the frequency or rate or breathing per minute (the total breathing frequency f_{TOT} is the sum of the mechanical f_{MECH} ventilator preselected frequency and the spontaneous f_{SPON} patient breathing frequency); the tidal volume (V_T), which is the volume of the breathing gas moving in and out of the lungs per breath ($V_{T MECH}$ is the ventilator preselected V_T per breath and $V_{T SPON}$ is the inhaled and exhaled volume per breath of the patient); the minute exhaled ventilation (V_E), which is the volume of breathing

25 gas moving in and out of the lungs of the patient per minute (V_E is the product of the breathing frequency f and the tidal volume ($V_E = f \times V_T$), and the $V_{E TOT}$ is the sum of the ventilator preselected V_E ($V_{E MECH}$) and the spontaneous patient V_E inhaled and exhaled per minute ($V_{E SPON}$)); the inhalation-to-exhalation time ratio (I:E ratio), which is the ratio of inhalation time to exhalation time during mechanical ventilation;

30

5 the physiologic dead space volume (V_{Dphys}), which is the volume of gas in the
anatomic airway and in ventilated, unperfused alveoli that does not participate in
blood gas exchange; the lung carbon dioxide elimination rate (LCO_2), which is the
volume of CO_2 exhaled per breath or per minute (LCO_2 is the area under the Ex CO_2
and volume curve); the partial pressure end-tidal carbon dioxide level ($PetCO_2$),
10 which is the partial pressure of the exhaled CO_2 measured at the end of the
exhalation; the cardiac output (CO) of the patient, which is the amount of blood
ejected from the heart per minute and which may, for example be derived from the
determined LCO_2 rate; the respiratory system compliance and resistance; the
respiratory muscle pressure, the work of breathing of the patient which may be
15 derived from the determined respiratory muscle pressure; and pressure-volume loops.

Ventilation data 72 may also be derived from the exemplified optional inputs
to the feature extraction subsystem 70. From the SPO2 output signal 51, the arterial
blood hemoglobin oxygen saturation level and the heart rate may be determined, and
20 the pulsatile blood pressure waveform of the SPO2 output signal 51 may be used to
determine arterial blood pressure. Additionally, from the blood pressure output signal
51, the arterial systolic, diastolic and mean blood pressure of the patient P may be
determined. Further, from the temperature output signal 51, the core body
temperature of the patient may 12 be derived. Still further, from the arterial blood
25 hemoglobin oxygen saturation level and the determined LCO_2 , the dead space volume
may be determined.

The feature extraction subsystem 70 may also receive user input via the
operator interface 60 and may receive the ventilator setting parameter signal 42. The
30 ventilation data 72 is preferably compiled in the feature extraction subsystem 70 and a
feature vector 74 or matrix is preferably generated which contains all of the
ventilation data items used by the monitor subsystem 10 to perform the ventilation
support assessment process. The feature vector 74 may be updated at user-defined
intervals such as, for example, after each breath or each minute and is output from the

5 feature extraction subsystem 70 to the intelligence subsystem 80 as a ventilation data output signal 75. Alternatively, as one skilled in the art will appreciate, the ventilation data 72 may be directly outputted to the intelligence subsystem 80 as the ventilation data output signal 75 without the intervening step of generating the feature vector 74 or matrix. The ventilation data 72 may also be outputted to the display 62.

10 Referring to Figs. 4, 6A and 6B, the intelligence subsystem 80 of the processing subsystem 40 preferably has a neural network 82. The primary function of the intelligence subsystem 80 is to make an assessment of the ventilator support provided to the patient and, based upon the assessment, recommend the desired level settings of the ventilator setting controls 30 which will adequately, and preferably optimally, support the physiological ventilation support needs of the patient P. For example, as shown in Fig. 6A, the intelligence subsystem 80 of the processing subsystem 40 may have a neural network 82 that receives the ventilation data output signal 75 containing the compiled ventilation data 72. The neural network 82 also receives the ventilator setting parameter signal 42 and may receive user input from the operator interface 60.

25 To fully appreciate the various aspects and benefits produced by the present invention, a basic understanding of neural network technology is required. Following is a brief discussion of this technology, as applicable to the ventilator monitor system 10 and method of the present invention.

30 Artificial neural networks loosely model the functioning of a biological neural network, such as the human brain. Accordingly, neural networks are typically implemented as computer simulations of a system of interconnected neurons. In particular, neural networks are hierarchical collections of interconnected processing elements configured, for example, as shown in FIG. 8. Specifically, FIG. 8 is a schematic diagram of a standard neural network 82 having an input layer 84 of processing elements, a hidden layer 86 of processing elements, and an output layer 88

5 of processing elements. The example shown in FIG. 8 is merely an illustrative embodiment of a neural network 82 that can be used in accordance with the present invention. Other embodiments of a neural network 82 can also be used, as discussed next.

10 Turning next to the structure of a neural network 82, each of its processing elements receives multiple input signals, or data values, that are processed to compute a single output. The output value is calculated using a mathematical equation, known in the art as an activation function or a transfer function that specifies the relationship between input data values. As known in the art, the activation function may include a
15 threshold, or a bias element. As shown in FIG. 8, the outputs of elements at lower network levels are provided as inputs to elements at higher levels. The highest level element, or elements, produces a final system output, or outputs.

20 In the context of the present invention, the neural network 82 is a computer simulation that is used to produce a recommendation of the desired ventilator setting of the ventilator controls 30 of the ventilator 20 which will adequately, and preferably optimally, support the physiological ventilation support needs of the patient, based upon at least a portion of the available ventilation setting parameters 42 and at least a portion of the ventilation data output signal 75 (*i.e.*, at least a portion of the derived
25 ventilation data 72).

30 The neural network 82 of the present invention may be constructed by specifying the number, arrangement, and connection of the processing elements which make up the network 82. A simple embodiment of a neural network 82 consists of a fully connected network of processing elements. The processing elements of the neural network 82 are grouped into layers: an input layer 84 where at least a portion of selected ventilation data 72, output signals 51, and the selected ventilator setting parameter signals 42 are introduced; a hidden layer 86 of processing elements; and an output layer 88 where the resulting determined level setting(s) for the control(s) 30 is

5 produced. The number of connections, and consequently the number of connection weights, is fixed by the number of elements in each layer.

10 In a preferred embodiment of the present invention, the data types provided at the input layer may remain constant. In addition, the same mathematical equation, or transfer function, is normally used by the elements at the middle and output layers. The number of elements in each layer is generally dependent on the particular application. As known in the art, the number of elements in each layer in turn determines the number of weights and the total storage needed to construct and apply the neural network 82. Clearly, more complex neural networks 82 generally require more configuration information and therefore more storage.

15 In addition to the structure illustrated in FIG. 6A, the present invention contemplates other types of neural network configurations for the neural network module such as the example shown in Fig. 6B, which is described in more detail below. All that is required by the present invention is that a neural network 82 be able to be trained and retrained, if necessary, for use to determine the desired level settings of the controls 30 of the ventilator 20. It is also preferred that the neural network 82 adapt (*i.e.*, learn) while in operation to refine the neural network's 82 determination of the appropriate level settings for the controls 30 of the ventilator 20.

25 Referring back to Figs. 6A and 8, the operation of a specific embodiment of a feedforward neural network 82 is described in more detail. It should be noted that the following description is only illustrative of the way in which a neural network 82 used in the present invention can function. Specifically, in operation, at least a portion of selected ventilation data 72 from the ventilation data output signal 75 and the selected ventilator setting parameter signals 42 (*i.e.*, collectively the input data) is provided to the input layer 84 of processing elements, referred to hereafter as inputs. The hidden layer elements are connected by links 87 to the inputs, each link 87 having an associated connection weight. The output values of the input processing elements

30

5 propagate along these links 87 to the hidden layer 86 elements. Each element in the
hidden layer 86 multiplies the input value along the link 87 by the associated weight
and sums these products over all of its links 87. The sum for an individual hidden
layer element is then modified according to the activation function of the element to
10 produce the output value for that element. In accordance with the different
embodiments of the present invention the processing of the hidden layer elements can
occur serially or in parallel.

15 If only one hidden layer 86 is present, the last step in the operation of the
neural network is to compute the output(s), or the determined level setting(s) of the
control(s) 30 of the ventilator by the output layer element(s). To this end, the output
values from each of the hidden layer processing elements are propagated along their
links 87 to the output layer element. Here, they are multiplied by the associated
weight for the link 87 and the products are summed over all links 87. The computed
20 sum for an individual output element is finally modified by the transfer function
equation of the output processing element. The result is the final output or outputs
which, in accordance with a preferred embodiment of the present invention, is the
desired level setting or settings of the ventilator setting controls 30.

25 In the example of the intelligence subsystem 80 shown in Fig. 6B, the
intelligence subsystem 80 is a hybrid intelligence subsystem that contains both rule-
based modules 90 as well as neural networks 82. In this alternative embodiment of
the intelligence subsystem 90, the determination of the desired level settings of the
controls 30 of the ventilator 20 are broken down into a number of tasks that follow
classical clinical paradigms. Each task may be accomplished using a rule-based
30 system 90 or a neural network 82. In the preferred configuration, the determination of
desired level settings of the ventilator setting controls 30 are performed by one of a
series of neural networks 82.

5 The purpose of the ventilation status module 92 is to make an initial
assessment of the adequacy of the ventilation support being provided to the patient P
based on the level settings of the ventilator setting controls 30 (as inputted to the
intelligence subsystem by the ventilator setting parameter signals 42) and the
ventilation data output signal. The final determination of the desired level settings of
10 the ventilator setting controls 30 is accomplished by one of a series of available neural
networks 82 in the ventilator control setting predictor module 94. The purpose of the
rule-based front end 96 is to determine, based on inputs from the ventilation status
module 92, data entered by the operator, and the ventilator setting parameter signal
42, which of the available neural networks 82 will determine the desired level settings
15 of the ventilator setting controls 30. The rule-based front end 96 will also determine
which inputs are extracted from the ventilation data output signal 75 and presented to
the selected neural network 82. Inputs to the ventilator control setting predictor
module 94 include ventilator data 72 from the ventilation data output signal 75, user
input, and input from the ventilator setting parameter signals 42. The purpose of the
20 rule-based back end module 98 is to organize information from previous modules,
neural networks 82, user input, and ventilation data 72 in the ventilation data output
signal and to format the information for display on the visual display 62 as well as for
storage to an external storage 64 such as a disk file.

25 As with most empirical modeling technologies, neural network development
requires a collection of data properly formatted for use. Specifically, as known in the
art, input data and/or the outputs of intermediate network processing layers may have
to be normalized prior to use. It is known to convert the data to be introduced into the
neural network 82 into a numerical expression, to transform each of the numerical
30 expressions into a number in a predetermined range, for example by numbers between
0 and 1. Thus, the intelligent subsystem of the present invention preferably has means
for: i) selecting at least a portion of the ventilation data 72 from the ventilation data
output signal 75 and at least a portion of the ventilator setting parameter signals 42, ii)
converting the selected portion of the ventilation data 72 and the selected portion of

5 the ventilator setting parameter signals 42 into numerical expressions, and iii)
transforming the numerical expressions into a number in a predetermined range.

10 In one conventional approach which can also be used in the present invention,
the neural network 82 of the present invention may include a preprocessor 83. The
preprocessor 83 extracts the correct data from the processing subsystem memory 48
and normalizes each variable to ensure that each input to the neural network 82 has a
value in a predetermined numerical range. Once the data has been extracted and
normalized, the neural network 82 is invoked. Data normalization and other
formatting procedures used in accordance with the present invention are known to
those skilled in the art and will not be discussed in any further detail.

15 In accordance with a preferred embodiment of the present invention the neural
network 82 is trained by being provided with the ventilator control setting assessment
made by a physician and with input data, such as ventilation data 72, the ventilation
control setting parameter signals 42, and the output signals 51 that were available to
the physician. In the sequel, the assessment along with the corresponding input
measurement and input data is referred to as a data record. All available data records,
possibly taken for a number of different patients, comprise a data set. In accordance
with the present invention, a data set corresponding is stored in memory and is made
25 available for use by the processing subsystem 40 for training and diagnostic
determinations.

30 A typical training mechanism used in a preferred embodiment of the present
invention is briefly described next. Generally, the specifics of the training process are
largely irrelevant for the operation of the ventilation monitor system. In fact, all that
is required is that the neural network 82 be able to be trained and retrained, if
necessary, such that it can be used to determine acceptably accurate determinations of
desired level settings of the controls 30 of the ventilator 20. Neural networks 82 are
normally trained ahead of time using data extracted from patients 12 by other means.

5 Using what it has learned from the training data, the neural network 82 may apply it to other/new patients P.

10 As known in the art, a myriad of techniques has been proposed in the past for training feedforward neural networks. Most currently used techniques are variations of the well-known error back-propagation method. The specifics of the method need not be considered in detail here. For further reference and more detail the reader is directed to the excellent discussion provided by Rumelhardt et al. in "Parallel Distributed Processing: Explorations in the Microstructure of Cognition," vols. 1 and 2, Cambridge: MIT Press (1986), and "Explorations in Parallel Distributed Processing, A Handbook of Models, Programs, and Exercises," which are
15 incorporated herein in their entirety by reference.

Briefly, in its most common form back-propagation learning is performed in three steps:

- 20
1. Forward pass;
 2. Error back-propagation; and
 3. Weight adjustment.

25 As to the forward pass step, in accordance with the present invention a single data record, which may be extracted from the ventilation data output signal 75 and the ventilator setting parameter signal(s) 42, is provided to the input layer 84 of the network 82. This input data propagates forward along the links 87 to the hidden layer elements which compute the weighted sums and transfer functions, as described above. Likewise, the outputs from the hidden layer elements are propagated along the
30 links to the output layer elements. The output layer elements computes the weighted sums and transfer function equations to produce the desired ventilator control settings 30.

5 In the following step of the training process, the physician assessment
associated with the data record is made available. At that step, the determination of
the desired level settings of the ventilator controls 30 produced by the neural network
82 is compared with the physician's assessment. Next, an error signal is computed as
the difference between the physician's assesment and the neural network's 82
10 determination. This error is propagated from the output element back to the
processing elements at the hidden layer 86 through a series of mathematical equations,
as known in the art. Thus, any error in the neural network output is partially assigned
to the processing elements that combined to produce it.

15 As described earlier, the outputs produced by the processing elements at the
hidden layer 86 and the output layer 88 are mathematical functions of their connection
weights. Errors in the outputs of these processing elements are attributable to errors
in the current values of the connection weights. Using the errors assigned at the
previous step, weight adjustments are made in the last step of the back-propagation
20 learning method according to mathematical equations to reduce or eliminate the error
in the neural network determination of the desired level setting of the ventilator
setting controls 30.

25 The steps of the forward pass, error back-propagation, and weight adjustment
are performed repeatedly over the records in the data set. Through such repetition, the
training of the neural network 82 is completed when the connection weights stabilize
to certain values that minimize, at least locally, the determination errors over the
entire data set. As one skilled in the art will appreciate however, the neural network
82 may, and preferably will, continue to train itself (*i.e.*, adapt itself) when placed into
30 operational use by using the data sets received and stored in the memory of the
processing subsystem 40 during operational use. This allows for a continual
refinement of the monitor 10 as it is continually learning, *i.e.*, training, while in
operational use. Further, it allows for the continual refinement of the determination of

5 the appropriate ventilator level settings in regard to the particular patient P to which
the ventilator 20 is operatively attached.

10 In addition to back-propagation training, weight adjustments can be made in
alternate embodiments of the present invention using different training mechanisms.
For example, as known in the art, the weight adjustments may be accumulated and
applied after all training records have been presented to the neural network 82. It
should be emphasized, however, that the present invention does not rely on a
15 particular training mechanism. Rather, the preferred requirement is that the resulting
neural network 82 produce acceptable error rates in its determination of the desired
level settings of the ventilator setting controls 30.

20 Upon completion of the determination of the desired level settings of the
ventilator setting controls 30 by the intelligent subsystem 80 of the processing system
40, the desired level settings of the ventilator setting controls 30 may be displayed on
the visual display 62 for use by the physician. The stored ventilation data output
signal 75, and particularly the subset of the ventilation data output signal 75
containing the ventilation data 72 that was used by the intelligent subsystem 80 in the
determination of the desired level setting of the controls 30, may be provided to the
25 visual display 62. Also, the stored ventilator setting parameter signals 42 and the
stored output signals 51 may be displayed on the visual display 62 in an appropriate
format. At this point, the physician can review the results to aid in her or his
assessment of the desirability of the recommended desired level settings of the
ventilator setting controls 30. The displayed results can be printed on printer [not
30 shown] to create a record of the patient's condition. In addition, with a specific
preferred embodiment of the present invention, the results can be communicated to
other physicians or system users of computers connected to the ventilator monitor
system 10 via an interface (not shown), such as for example a modem or other method
of electronic communication.

5 Additionally, a preferred embodiment the present invention provides a
real-time ventilator monitor system 10 and method. Real-time operation demands, in
general, that input data be entered, processed, and displayed fast enough to provide
immediate feedback to the physician in the clinical setting. In alternate embodiments,
off-line data processing methods can be used as well. In a typical off-line operation,
10 no attempt is made to respond immediately to the physician. The measurement and
interview data in such case is generated some time in the past and stored for retrieval
and processing by the physician at an appropriate time. It should be understood that
while the preferred embodiment of the present invention uses a real-time approach,
alternative embodiments can substitute off-line approaches in various steps.

15 The preferred method of operation of the present invention comprises the steps
of receiving at least one ventilator setting parameter signal 42 indicative of the current
level settings of the controls 30 of the ventilator 20, monitoring a plurality of output
signals 51 to determine the sufficiency of ventilation support supplied to the patient P,
20 determining the desired level settings of the ventilator setting controls 30, and
displaying the desired level settings of the controls 30 to the operating clinician.

25 The output signals 51 received may comprise a plurality of signals selected
from a group of: an exhaled carbon dioxide signal indicative of the exhaled carbon
dioxide (ExCO₂) level of the exhaled gas expired by the patient P within the
breathing circuit 22; a flow rate signal indicative of the flow rate (V) of the
inhaled/exhaled gas expired by patient P within the breathing circuit 22; a pulse
oximeter hemoglobin oxygen saturation (SpO₂) signal indicative of the oxygen
saturation level of the patient P; a pressure (P) signal indicative of the pressure of the
30 breathing gas within the breathing circuit 22; a blood pressure (BP) signal indicative
of the blood pressure of the patient 12. The output signals 51 may also comprise a
temperature (T) signal indicative of the core body temperature of the patient P, an
arterial blood gas PaO₂ signal, an arterial blood gas PaCO₂ signal, and/or an arterial
blood gas pH signal.

5 The ventilator setting parameter signal 42 may comprise at least one of: a
minute ventilation (V_E) signal indicative of the V_E level set on the ventilator 20; a
ventilator breathing frequency (f) signal indicative of the f level set on the ventilator
20; a tidal volume (V_T) signal indicative of the V_T level set on the ventilator 20; a
breathing gas flow rate (V) signal indicative of the V level set on the ventilator 20; a
10 pressure limit signal indicative of the pressure limit set on the ventilator 20; a work of
breathing (WOB) signal indicative of the WOB level set on the ventilator 20; a
pressure support ventilation (PSV) signal indicative of the PSV level set on the
ventilator 20; a positive end expiratory pressure (PEEP) signal indicative of the PEEP
15 level set on the ventilator 20; a continuous positive airway pressure (CPAP) signal
indicative of the CPAP level set on the ventilator 20; and a fractional inhaled oxygen
concentration (FIO2) signal indicative of the FIO2 level set on the ventilator 20.

20 For example, the step of determining the desired level settings of the ventilator
setting controls 30 of the ventilator 20 may comprise the steps of generating
ventilation data 72 from the received output signals 51 in the processing subsystem 40
and applying at least a portion of the generated ventilation data 72 and the ventilator
setting parameter signal 42 to the neural network 82 of the processing subsystem 40.
If desired, at least a portion of the output signals 51 may also be applied to the neural
25 network 82 as ventilation data 72.

30 In an alternative example, the step of determining the desired level settings of
the controls 30 of the ventilator 20 may comprise the steps of generating ventilation
data 72 from the received output signals 51 in the processing subsystem 40, applying
a set of decision rules in the rule based front-end 96 to at least a portion of the
ventilation data 72 and the ventilator setting parameter signal 42 to classify the
applied portions of the ventilation data 72 and the ventilator setting parameter signal
42, selecting an appropriate neural network 82 to use, and applying a portion of the
ventilation data 72 and the ventilator setting parameter signal 42 to the selected neural

5 network 82 which will be used to determine the desired level settings of the ventilator setting controls 30.

10 The ventilator monitor system 10 of the present invention may be implemented in one of many different configurations. For example, the ventilator monitor system 10 may be incorporated within a ventilator 20. In an alternative example, the ventilator monitor system 10 may be a stand alone monitor that is operatively connected to the ventilator 20.

15 A realization of an embodiment of the processing subsystem 40 of the present invention is illustrated in Fig. 7. Here, the processing subsystem 40 includes the processor 46, which is preferably a microprocessor, memory 48, storage devices 64, controllers 45 to drive the display 62, storage 64, and ventilator 20, and an analog-to-digital converter (ADC) 47 if required. The processing subsystem 40 also includes a neural network 82, which may, for example, be embodied in a neural network board 20 49. The ADC and neural network boards 47, 49 are commercially available products. There is also an optional output board (not shown) for connection to a computer network and/or central monitoring station.

25 The ADC board 47 converts the analog signal received from the output of any of the sensors 52 of the measuring means to a digital output that can be manipulated by the processor 46. In an alternative implementation, the output of any of the sensors 52 could be connected to the processor 46 via digital outputs, e.g., a serial RS232 port. The particular implementation is determined by the output features of the particular sensor 52. The processor 46 should contain circuits to be programmed 30 for performing mathematical functions, such as, for example, waveform averaging, amplification, linearization, signal rejection, differentiation, integration, addition, subtraction, division and multiplication, where desired. The processor 46 may also contain circuits to be programmed for performing neural/intelligent control software, neural network learning software, and ventilator control software, as required.

5 Circuits or programs performing these functions are well known to one skilled in the art, and they form no part of the present invention. The processor 46 executes the software which makes the computations, controls the ADC and neural network boards 47, 49, and controls output to the display and storage devices 62, 64, network communication, and the ventilator apparatus 20.

10 The purpose of the neural network board 49 is to implement the neural/intelligent control software. As one skilled in the art will appreciate, the need for a separate neural network board 49 is determined by the computational power of the main processor 46. With recent increases in microprocessor speeds, it may not be necessary to have a separate board 49, since some or all of these functions could be handled by the processor 46. The need for the separate board 49 is also determined by the precise platform on which the invention is implemented.

15 In addition, while the processor 46 of the processing subsystem 40 has been described as a single microprocessor, it should be understood that two or more microprocessors could be used dedicated to the individual functions. For example, the ventilator 20 may have a microprocessor that is operatively coupled to the processing subsystem 40 of the monitor system 10. In this manner the monitor system 20 could be incorporated into a modular system 10 that may be coupled to any conventional microprocessor-controlled ventilator 20 for monitoring of the ventilation support provided by the ventilator 20. Alternatively, as one skilled in the art will appreciate, and as shown in Fig. 2B, the monitor system 10 of the present invention may be incorporated into the design of a microprocessor-controlled ventilator 10 with the processing subsystem 40 of the ventilator monitor system using the microprocessor of the ventilator 20. In addition, the functions of the processor 46 could be achieved by other circuits, such as application specific integrated circuits (ASIC), digital logic circuits, a microcontroller, or a digital signal processor.

5 The invention has been described herein in considerable detail, in order to
comply with the Patent Statutes and to provide those skilled in the art with
information needed to apply the novel principles, and to construct and use such
specialized components as are required. However, it is to be understood that the
invention can be carried out by specifically different equipment and devices, and that
10 various modification, both as to equipment details and operating procedures can be
effected without departing from the scope of the invention itself. Further, it should be
understood that, although the present invention has been described with reference to
specific details of certain embodiments thereof, it is not intended that such details
should be regarded as limitations upon the scope of the invention except as and to the
15 extent that they are included in the accompanying claims.

What is claimed is:

1. A ventilation support monitoring method for a ventilator supplying a breathing gas to a patient via a breathing circuit in fluid communication with at least one lung of the patient, the ventilator having a plurality of selectable ventilator setting controls for governing supply of the breathing gas from the ventilator to the patient, each setting control selectable to a level setting, comprising:

receiving at least one ventilator setting parameter signal, each ventilator setting parameter signal indicative of the level setting of one ventilator setting control;

monitoring a plurality of sensors to determine the sufficiency of ventilation support supplied to the patient, each sensor operatively connected to a select one of the patient or the breathing circuit, each sensor generating an output signal; and

determining the desired level setting of at least one ventilator setting control of the ventilator.

2. The method of Claim 1, wherein the output signals are selected from the group comprising: an exhaled carbon dioxide signal indicative of the exhaled carbon dioxide (ExCO₂) level of the exhaled gas expired by the patient within the breathing circuit; a flow rate signal indicative of the flow rate (V) of the inhaled/exhaled gas expired by patient within the breathing circuit; a pulse oximeter hemoglobin oxygen saturation (SpO₂) signal indicative of the oxygen saturation level of the patient; a pressure (P) signal indicative of the pressure of the breathing gas within the breathing circuit; a blood pressure (BP) signal indicative of the blood pressure of the patient; and a temperature (T) signal indicative of the core body temperature of the patient.

3. The method of Claim 2, wherein the output signals also comprise at least one of: an arterial blood gas PaO₂ signal; an arterial blood gas PaCO₂ signal; and an arterial blood gas pH signal.

4. The method of Claim 1, wherein the ventilator setting parameter signal comprises at least one of: a minute ventilation (V_E) signal indicative of the V_E level set on the ventilator; a ventilator breathing frequency (f) signal indicative of the f level set on the ventilator; a tidal volume (V_T) signal indicative of the V_T level set on the ventilator; a breathing gas flow rate (V) signal indicative of the V level set on the ventilator; a pressure limit signal indicative of the pressure limit set on the ventilator; a work of breathing (WOB) signal indicative of the WOB level set on the ventilator; a pressure support ventilation (PSV) signal indicative of the PSV level set on the ventilator; a positive end expiratory pressure (PEEP) signal indicative of the PEEP level set on the ventilator; a continuous positive airway pressure (CPAP) signal indicative of the CPAP level set on the ventilator; and a fractional inhaled oxygen concentration (FIO_2) signal indicative of the FIO_2 level set on the ventilator.

5. The method of Claim 1, wherein monitoring the sensors comprises measuring a plurality of ventilation support parameters with the plurality of sensors, each sensor connected to a processing subsystem, wherein each sensor generates one output signal corresponding to the particular measured ventilation support parameter.

6. The method of Claim 1, further comprising displaying the desired level settings of the ventilator setting controls.

7. The method of Claim 1, wherein determining the desired level settings of the ventilator setting controls further comprises generating ventilation data from the output signals in a processing subsystem.

8. The method of Claim 7, wherein the processing subsystem has a neural network, and wherein determining the desired level settings of the ventilator setting controls of the ventilator further comprises applying at least a portion of the ventilation data and the ventilator setting parameter signal to the neural network of the

processing subsystem to determine the desired level settings of the ventilator setting controls.

9. The method of Claim 7, wherein the processing subsystem has a neural network, and wherein the step of determining the desired level settings of the ventilator setting controls further comprises:

applying a set of decision rules to at least a portion of the ventilation data and the ventilator setting parameter signal to classify a portion of the ventilation data and the ventilator setting parameter signal;

applying a portion of the ventilation data and the ventilator setting parameter signal to the neural network; and

determining at least one desired level settings of the ventilator setting controls of the ventilator from the applied portion of the ventilation data and the ventilator setting parameter signal.

10. A method for differential determination of a plurality of desired level settings of a ventilator having selectable ventilator setting controls, each setting control selectable to a level setting, the method comprising:

supplying a breathing gas from the ventilator to a patient via a breathing circuit in fluid communication with the ventilator and at least one lung of the patient;

receiving a plurality of output signals indicative of the physiological characteristics of the patient and the characteristics of the breathing gas supplied to the patient;

receiving a plurality of ventilator setting parameter signals indicative of the level settings of the ventilator setting controls;

deriving a plurality of ventilation data from the output signals;

selecting at least a portion of the ventilation data and at least a portion of the ventilator setting parameter signals;

converting the selected portion of the ventilation data and the selected portion of the ventilator setting parameter signals into numerical expressions;

transforming each of the numerical expressions into a number in a predetermined range;

inputting a plurality of the transformed numerical expressions into a neural network; and

determining at least one of the desired level settings of the ventilator setting controls using the neural network in accordance with the input numerical expressions.

11. The method of Claim 10, further comprising training the neural network to determine each of the desired level settings of the ventilator setting controls using the ventilation data and the ventilator setting parameter signals.

12. The method of Claim 11, further comprising dividing the determination of the desired level settings of the ventilator setting controls into a plurality of stages using a plurality of neural networks.

13. The method of Claim 11, wherein the neural network includes a plurality of parallel neural networks, each of the parallel neural networks having a plurality of inputs and one output, the outputs respectively corresponding to the plurality of setting controls, so that one output corresponds to one desired level setting of one ventilator setting control.

14. The method of Claim 10, wherein deriving the ventilation data comprises deriving a plurality of: pressure (P) of the breathing gas within the breathing circuit; flow rate (V) of the breathing gas within the breathing circuit; exhaled carbon dioxide (ExCO₂) in the breathing gas within the breathing circuit; peak inflation pressure (PIP); mean airway pressure (Paw); positive end expiratory pressure (PEEP); continuous positive airway pressure (CPAP); breathing frequency (f); tidal volume (V_T); minute exhaled ventilation (V_E); inhalation-to-exhalation time ratio (I:E); physiological dead space volume (V_{dphys}); lung carbon dioxide elimination rate (LCO₂); partial pressure end-tidal carbon dioxide (PetCO₂); respiratory muscle

pressure (RMP) of the patient; work of breathing (WOB) of the patient; cardiac output (CO) of the patient; pulse oximeter hemoglobin oxygen saturation (SpO₂) of the patient; heart rate (HR) of the patient; blood pressure (BP) of the patient; arterial blood gas PaO₂ of the patient; arterial blood gas PaCO₂ of the patient, arterial blood gas pH of the patient; and temperature (T) of the patient.

15. The method of Claim 10, wherein receiving the ventilator setting parameter signals comprises receiving a plurality of signals selected from signals indicative of: a minute ventilation (V_E) level set on the ventilator; a ventilator breathing frequency (f) level set on the ventilator; a tidal volume (V_T) level set on the ventilator; a breathing gas flow rate (V) level set on the ventilator; a pressure limit level set on the ventilator; a work of breathing (WOB) level set on the ventilator; a pressure support ventilation (PSV) level set on the ventilator; a positive end expiratory pressure (PEEP) level set on the ventilator; a continuous positive airway pressure (CPAP) level set on the ventilator; and a fractional inhaled oxygen concentration (FIO₂) level set on the ventilator.

16. The method of Claim 10, wherein the neural network comprises a plurality of input units, a plurality of hidden layers having a plurality of hidden units, and a plurality of output units, the output units respectively corresponding to the plurality of ventilator settings.

17. The method of Claim 10, further comprising:
selecting a subset of the output signals for display; and
displaying the selected subset of output signals in real time.

18. The method of Claim 10, further comprising displaying at least one of the plurality of desired level settings for the ventilator setting controls of the ventilator.

19. A ventilation support monitoring system for a ventilator supplying a breathing gas to a patient via a breathing circuit in fluid communication with at least one lung of a patient, the ventilator having a plurality of selectable ventilator setting controls for governing supply of the breathing gas from the ventilator to the patient, each setting control selectable to a level setting, and each ventilator setting control generating a ventilator setting parameter signal indicative of the current level setting of the ventilator setting control, the system comprising:

an input that receives at least one ventilator setting parameter signal;

a plurality of sensors for measuring a plurality of ventilation support parameters, each sensor operatively connecting to a select one of the patient or the breathing circuit, wherein each sensor generates an output signal based on the measured ventilation support parameter; and

a processing subsystem connected to receive the output signals from the sensors and the ventilator setting parameter signal from the input, the processing subsystem having a processor and a memory, the processor running under control of a program stored in the memory, the processing subsystem having an intelligence system to determine a desired level setting of at least one ventilator setting control in response to the ventilator setting parameter signal and the output signals.

20. The system of Claim 19, wherein the plurality of ventilation support parameters is selected from the group comprising: the flow rate (V) of the exhaled gas inspired/expired by patient within the breathing circuit; the exhaled carbon dioxide (ExCO₂) level of the exhaled gas expired by the patient within the breathing circuit; the hemoglobin oxygen saturation (SpO₂) level of the patient; the pressure (P) of the breathing gas within the breathing circuit; the blood pressure (BP) of the patient; and the core body temperature (T) of the patient.

21. The system of Claim 20, wherein the means the plurality of ventilation support parameters also includes at least one of: the arterial blood gas PaO₂ level of the

patient; the arterial blood gas PaCO₂ level of the patient; and the arterial blood gas pH level of the patient.

22. The system of Claim 19, wherein the ventilator setting parameter signal comprises at least one of: a minute ventilation (V_E) signal indicative of the V_E level set on the ventilator; a ventilator breathing frequency (f) signal indicative of the f level set on the ventilator; a tidal volume (V_T) signal indicative of the V_T level set on the ventilator; a breathing gas flow rate (V) signal indicative of the V level set on the ventilator; a pressure limit signal indicative of the pressure limit set on the ventilator; a work of breathing (WOB) signal indicative of the WOB level set on the ventilator; a pressure support ventilation (PSV) signal indicative of the PSV level set on the ventilator; a positive end expiratory pressure (PEEP) signal indicative of the PEEP level set on the ventilator; a continuous positive airway pressure (CPAP) signal indicative of the CPAP level set on the ventilator; and a fractional inhaled oxygen concentration (FIO₂) signal indicative of the FIO₂ level set on the ventilator.

23. The system of Claim 19, further comprising a display, wherein the processing subsystem provides the desired level settings of the ventilator setting controls to the display.

24. The ventilator of Claim 19, further comprising an alarm for notifying an operator of the ventilator that the level settings of the ventilator setting controls differs from the determined desired level settings of the ventilator setting controls.

25. The system of Claim 19, wherein the desired level setting for the ventilator setting control of the ventilator comprises at least one of: a minute ventilation (V_E) level indicative of the desired V_E level to set on the ventilator; a ventilator breathing frequency (f) level indicative of the desired f level to set on the ventilator; a tidal volume (V_T) level indicative of the V_T level to set on the ventilator; a breathing gas flow rate (V) level indicative of the V level to set on the ventilator; a pressure limit

level indicative of the pressure limit level to set on the ventilator; a work of breathing (WOB) level indicative of the WOB level to set on the ventilator; a pressure support ventilation (PSV) level indicative of the PSV level to set on the ventilator; a positive end expiratory pressure (PEEP) level indicative of the PEEP level to set on the ventilator; a continuous positive airway pressure (CPAP) level indicative of the CPAP level to set on the ventilator; and a fractional inhaled oxygen concentration (FIO2) level indicative of the FIO2 level to set on the ventilator.

26. The system of Claim 23, wherein:

the processing subsystem has at least one neural network ; and
the processor, in determining the desired level settings of the ventilator setting controls, generates a plurality of ventilation data from the output signals of the sensors and applies at least a portion of the ventilation data and at least a portion of the ventilator setting parameter signal to the neural network to generate the desired level settings of the ventilator setting controls.

27. The system of Claim 26, wherein the processing subsystem is programmed with a set of decision rules and wherein the processor applies the set of decision rules to the ventilation data prior to applying the portion of the ventilation data and the portion of the ventilator setting parameter signal to the neural network.

28. The system of Claim 26, further comprising a display, wherein the processing subsystem identifies ventilation data used to determine the desired level settings of the ventilator setting controls, identifies a subset of the ventilation data for display, and provides the subset of the ventilation data to the display, and wherein the processing subsystem provides the desired level settings of the ventilator setting controls to the display.

29. The system of Claim 23, wherein the processing subsystem has at least one neural network, the neural network under control of a program stored in the memory, and wherein the determining means of the processing subsystem comprises:

means for generating ventilation data from the received output signals;

means for selecting at least a portion of the ventilation data and at least a portion of the ventilator setting parameter signals;

means for converting the selected ventilation data and the selected ventilator setting parameter signal into numerical expressions;

means for transforming each of the numerical expressions into a number in a predetermined range;

means for inputting a plurality of the transformed numerical expression into the neural network so that the desired level settings of the ventilator setting controls are determined in accordance with the input numerical expressions.

30. The system of Claim 23, wherein the processing subsystem has at least one neural network, the neural network under control of a program stored in the memory, and wherein the determining means of the processing subsystem has:

means for generating a plurality of training data sets, each training data set including output signals and indicated level settings of the ventilator setting controls associated with an historical occurrence of physiologic condition of the patient during ventilation support;

means for identifying the statistically significant training data sets of the plurality of training data sets; and

means for training the neural network using the statistically significant training data sets so that the desired level settings of the ventilator setting controls are determined based upon selected output signals and selected level settings of the ventilator setting controls.

31. A ventilation support monitoring system for a ventilator supplying a breathing gas to a patient via a breathing circuit in fluid communication with at least one lung of

a patient, the ventilator having a plurality of ventilator setting controls governing supply of the breathing gas from the ventilator to the patient, each setting control selectable to a level setting, and each ventilator setting control generating a ventilator setting parameter signal indicative of the current level setting of the ventilator setting control, the system comprising:

- an input that receives at least one ventilator setting parameter signal;
- a plurality of sensors for measuring a plurality of ventilation support parameters, each sensor generating an output signal indicative of the measured ventilation support parameter; and
- a processing subsystem having:
 - a neural network that receives data,
 - a processor having a memory, the processor connected to receive output signals from the sensors and the ventilator setting parameter signals from the input and running under control of a program stored in the memory to generate a plurality of ventilation data from the output signals, to apply at least a portion of the ventilation data and at least a portion of the ventilator setting parameter signals to the neural network to determine the desired level settings of the ventilator setting controls.

32. The system of Claim 31, wherein the plurality of ventilation support parameters comprises one or more of: the flow rate (V) of the exhaled gas inspired/expired by patient within the breathing circuit; the exhaled carbon dioxide (ExCO₂) level of the exhaled gas expired by the patient within the breathing circuit; the hemoglobin oxygen saturation (SpO₂) level of the patient; the pressure of the breathing gas within the breathing circuit; the blood pressure (BP) of the patient; and the core body temperature (T) of the patient.

33. The system of Claim 34, further comprising a display, wherein the processing subsystem provides the desired level settings of the ventilator setting controls to the display.

34. The system of Claim 31, wherein the desired level settings for the ventilator setting controls include at least one of the group comprising: a minute ventilation (V_E) level indicative of the desired V_E level to set on the ventilator; a ventilator breathing frequency (f) level indicative of the desired f level to set on the ventilator; a tidal volume (V_T) level indicative of the V_T level to set on the ventilator; a breathing gas flow rate (V) level indicative of the V level to set on the ventilator; a pressure limit level indicative of the pressure limit level to set on the ventilator; a work of breathing (WOB) level indicative of the WOB level to set on the ventilator; a pressure support ventilation (PSV) level indicative of the PSV level to set on the ventilator; a positive end expiratory pressure (PEEP) level indicative of the PEEP level to set on the ventilator; a continuous positive airway pressure (CPAP) level indicative of the CPAP level to set on the ventilator; and a fractional inhaled oxygen concentration (FIO_2) level indicative of the FIO_2 level to set on the ventilator.

35. The system of Claim 31, wherein the processor is programmed with a set of decision rules and wherein the processor applies the set of decision rules to the ventilation data prior to applying the portion of the ventilation data and the portion of the ventilator setting parameter signal to the neural network.

36. The system of Claim 33, wherein the processing subsystem identifies ventilation data used to determine the desired level settings of the ventilator setting controls; identifies a subset of the ventilation data for display; and provides the subset of the ventilation data to the display.

38. The ventilator of Claim 31, further comprising:
an alarm signal produced by the processing subsystem when at least one level setting
of the ventilator setting control differs from the desired level setting for that ventilator
setting control; and
one or more of an audible and visible alert in response to the alarm signal to alert the
operator.

[illegible]

Embodiments of the present invention described and shown in the specification and drawings include an system and method for monitoring the ventilation support provided by a ventilator that is supplying a breathing gas to a patient via a breathing circuit that is in fluid communication with the lungs of the patient. The ventilator has a plurality of selectable ventilator setting controls governing the supply of ventilation support from the ventilator, each setting control selectable to a level setting. The ventilator support monitor system preferably receives at least one ventilator setting parameter signal, each ventilator setting parameter signal indicative of the level settings of one ventilator setting control, monitors a plurality of sensors, each sensor producing an output signal indicative of a measured ventilation support parameter, to determine the sufficiency of the ventilation support received by the patient, and determines the desired level settings of the ventilator setting controls in response to the received ventilator setting parameter signal and the output signals. The ventilator support monitor system preferably utilizes a trainable neural network to determine the desired level settings of the ventilator setting controls. It is emphasized that this abstract is provided to comply with the rules requiring an abstract which will allow a searcher or other reader to quickly ascertain the subject matter of the technical disclosure. It is submitted with the understanding that it will not be used to interpret or limit the scope or meaning of the claims. 37 C.F.R. § 1.72(b).

Fig. 2A

10

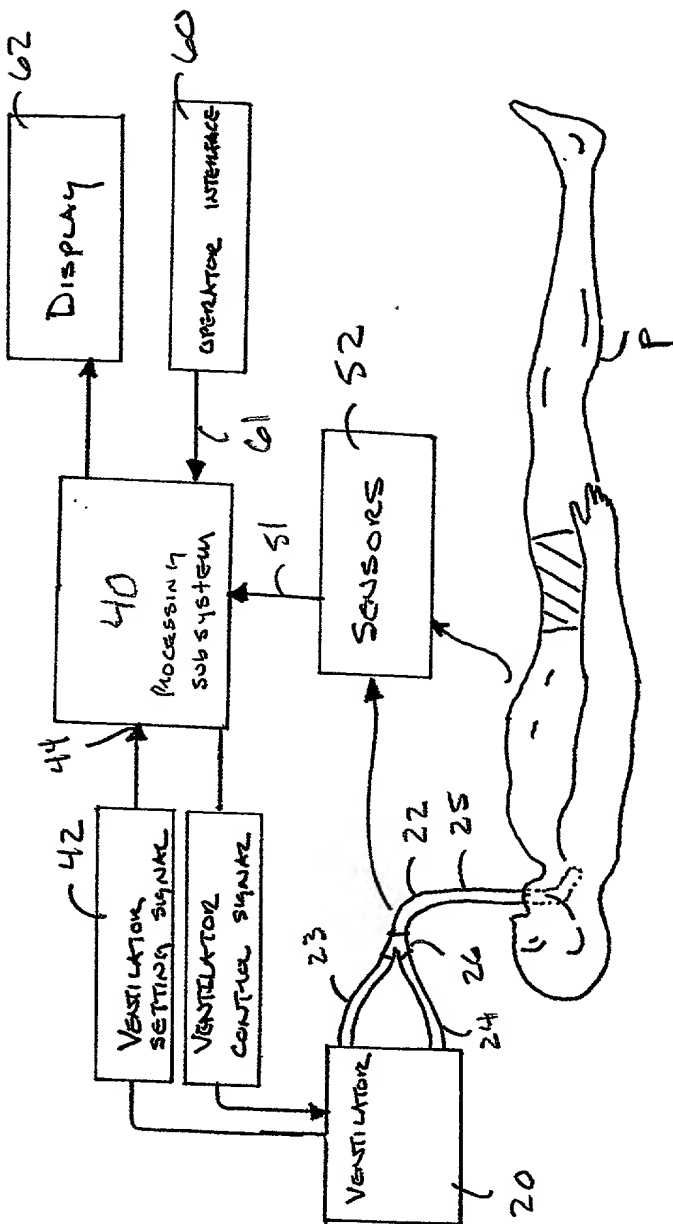


Fig. 2B

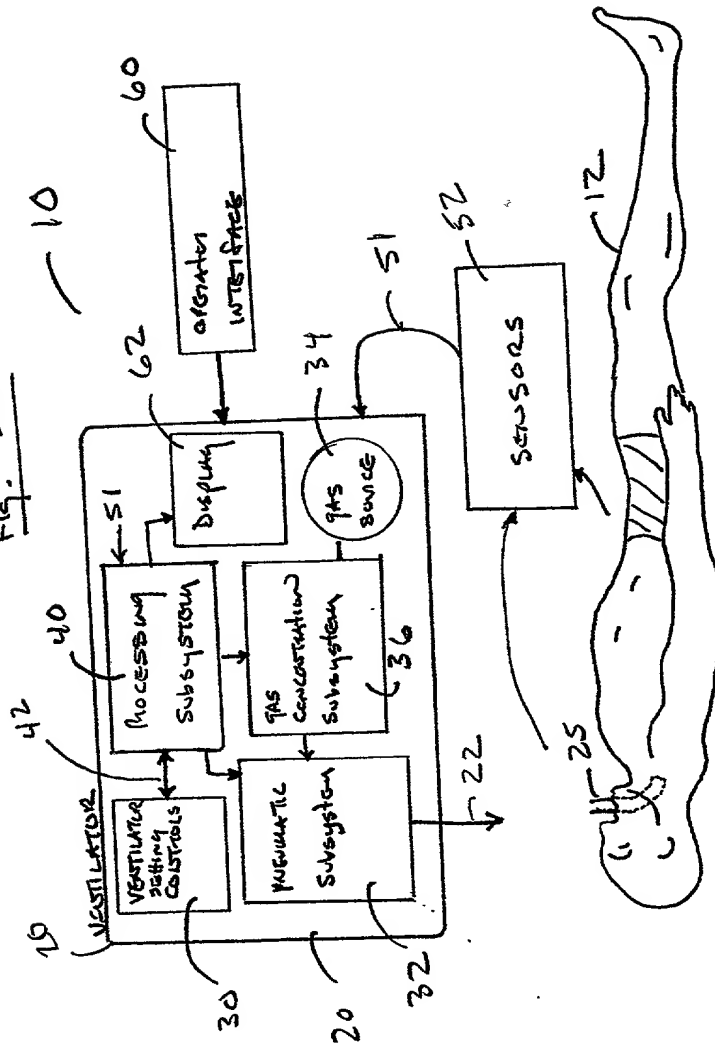
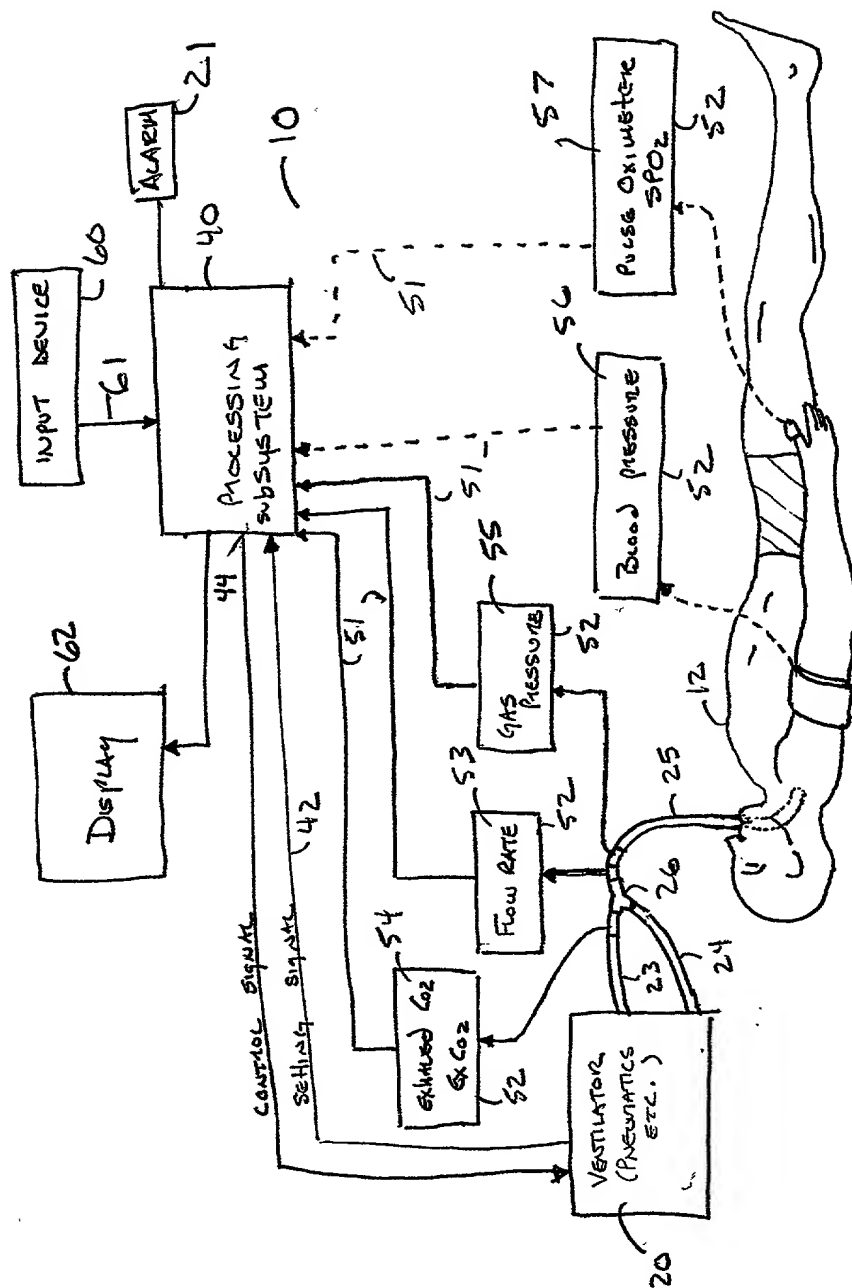


Fig. 3



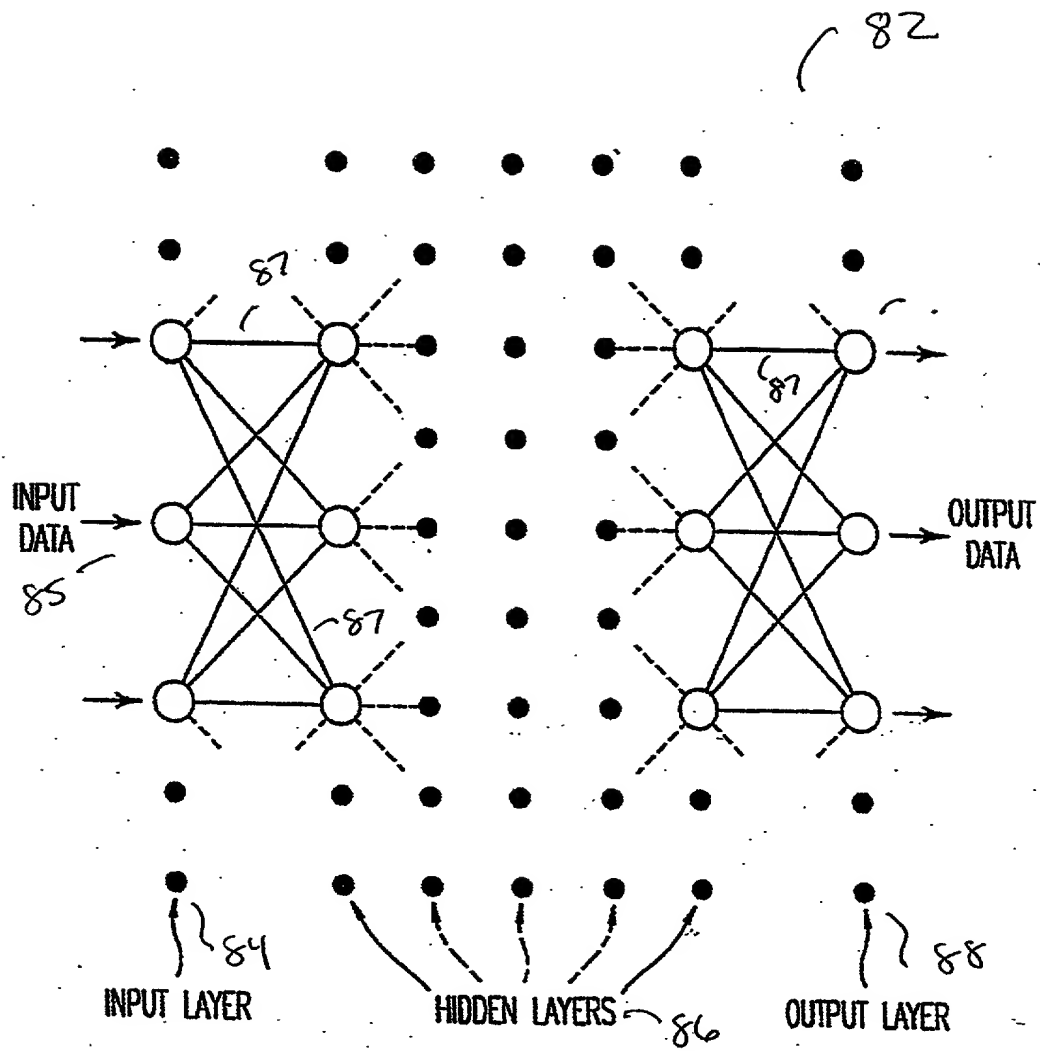


FIG. 8